

DOCKET NO.: CXU-363

**Fluoropolymer Compositions, Optical Devices, and Methods for
Fabricating Optical Devices**

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Title of the Invention

- 5 Fluoropolymer Compositions, Optical Devices, and Methods for
Fabricating Optical Devices

Cross Reference to Related Applications

- 10 This application is related to U.S. Serial No. 09/604,748, which is a
regular U.S. utility patent application filed 27 June 2000.

Field of the Invention

- This invention relates to the use of fluoropolymers and methods of
applying fluoropolymers in making components for optical applications. In
15 particular, the invention relates to perfluorocyclobutyl-based compositions.

Background of the Invention

- Optical waveguides interconnect optical information processing devices,
or connect such devices with other optical communication links such as glass
optical fibers. Waveguides may be used to create passive optical devices such as
20 splitters, combiners, couplers, routers and the like. In commonly used planar
applications, waveguides are densely packed on substrates.

- An optical waveguide typically comprises a transparent core that is
capable of directing light signals therethrough, and a cladding comprising a
material that affords a lower refractive index than the core material.
25 Waveguides may be constructed as single monolithic structures lithographically
provided on a substrate, or may comprise optical fibers.

- In the electronics and optical fabrication technologies, optical
interconnects have been used in backplane interconnections, board-to-board
interconnections, clock distribution, and a variety of other applications. In
30 particular, lithographic processes have been used because such processes are
generally suitable for mass production.

The integration of polymeric materials in optics is an increasingly attractive alternative in devices such as switches, optical interconnects, splitters, and surface relief structures. As demand for band width and low cost integration has increased, polymers provide flexibility, high transparency and versatility in structure, and properties. In particular, fluoropolymers represent alternatives to current optical polymers due to their properties, such as low transmission loss (at 1300 and 1550 nm), low birefringence, good optical stability after thermal aging, and low moisture absorption. For example, fluoroacrylates developed by Allied Signal and others, Dupont's Teflon™ AF (tetrafluoroethylene and perfluorovinyl ether copolymer), Amoco's Ultradel™ (fluorinated polyimide) and Asahi's CYTOP™ (perfluorovinyl ether cyclopolymer) are fluoroplastics currently pursued for optical device manufacture. See, for example, Eldad, L.; Schacklette, L. "Advances in Polymer Integrated Optics," *IEEE J. Quantum Electronics* 2000, 6(1), 54.

The use of a perfluorocyclobutyl-based homopolymer in an optical waveguide has been disclosed. See Fishbeck, G.; Moosbuerger, R.; Kostrzews, C.; Achen, A.; Petermann, K. *Electronic Letters* 1997, 33(6), 518. Also, the use of various perfluorocyclobutane (PFCB) homopolymers has been disclosed for various electronics and other applications. See for example, United States Patent No. 5,159,038; U.S. Patent No. 5,037,917 and related patents.

Several patents relate to optical devices and methods for constructing them. For example, United States Patent No. 5,850,498 is directed to low stress optical waveguides. United States Patent No. 6,210,867 B1 is directed to methods for fabricating low loss optical devices using a photoresist coated on a linear optical layer by a spin coating method. The photoresist is dipped into a developer fluid and baked, thereby forming a photoresist pattern defining specific areas upon which a metal layer is deposited. A vacuum deposition method such as sputtering, electron beam or thermal evaporation may be employed to deposit a metal substrate.

A pending United States patent application (U.S.S.N. 09/604,748) which is commonly owned by the assignee of the present application, discloses optical fluropolymers and methods of applying fluoropolymers in molding processes. The application is directed to the use of alternating perfluorocyclobutane and 5 aryl ether linkages that are adapted for micromolding polymeric films by replicating a pattern or image directly from a silicon master.

In the case of forming a polymer film using spin coating techniques, it is necessary to form a coating of sufficient thickness to manufacture a waveguide. Many polymers cannot be dissolved at solids content when spin coated to 10 manufacture a suitable waveguide thickness. In many instances, it requires multiple coats of polymer to achieve the necessary thickness, which introduces added interface problems, and can be costly and time consuming.

It has been recognized in the industry that a need exists for a suitable polymeric material and process that affords appropriate optical properties when 15 applied in an optical device. A polymer and process of applying the polymer that is capable of achieving a suitably thick coating on the device is needed. A polymeric composition that is capable of maintaining a suitable solids thickness in solvent, thereby affording a relatively thick coating, is desirable. A composition and method of application that employs a minimum amount of 20 solvent, and a maximum solids content, is needed. Furthermore, polymeric compositions that exhibits low loss in the telecommunications wavelength and a compositionally controlled refractive index to match that of the silica optical fibers and other components would be desirable.

Summary of the Invention

25 A method and solution for making an optical device is provided in the invention of the application. The method may include steps of: providing a perfluorocyclobutyl-based copolymer composition, coating the perfluorocyclobutyl-based copolymer composition upon a substrate to form a first film, and thermally curing the first film to form a thermoset film.

The invention comprises compositions that include fluoropolymers comprised of alternating perfluorocyclobutane and aryl ether linkages. These compositions possess the mechanical, thermal and structural properties and the desired optical clarity required for the manufacture of optical waveguides and
5 the like. Chemical structures of some of the compositions that may be employed in the invention of the application are shown in the attached drawings.

In some applications of the invention, the fluoropolymer may be used and as an optical signal carrier in a structural polymer matrix composite. Such composites may provide fully integrated high-speed data transmission mediums
10 with switching and routing capabilities. Thermoplastic and thermosetting polymers containing perfluorocyclobutane (PFCB) and arylene ether linkages provide high performance and a multi-functional polymer. PFCB segments may provide improved dielectric insulation, optical clarity in the visible and near infrared wavelengths, increased solubility, melt processability, large thermo-
15 optic coefficient (-dn/dT), low birefringence, and other valuable properties.

Of course, the above groups may be substituted with other alkyl or haloalkyl or organic constituents on the one or more carbons, such as a carbon in the ring structure, and the above structures are provided by way of example and not by way of limitation. In some applications, a polymer is provided which is
20 prepared from a trifluorovinyl aromatic ether (TVE). Trifluorovinyl aromatic ether monomer precursors are traditionally prepared in two high yielding steps from commercially available phenolic precursors such as, for example, tris (hydroxyphenyl) ethane, biphenol, bis(hydroxyphenyl) fluorene, or other compounds. Partially inorganic monomers containing siloxane and phosphine
25 oxide groups have also been synthesized using an intermediate strategy involving delivery of a TVE group intact via Grignard and organo-lithium chemistry.

Polymers with a wide range of mechanical, thermal and optical properties may be obtained using the above referenced chemistry. In particular,
30 the refractive indices of the homopolymers or the co-polymers may be provided

exactly as designed by careful regulation of monomer choice. Thus the refractive index at a given wavelength can be precisely controlled. The invention is capable of providing well defined cyclopolymerization mechanisms using PFCB to prepare polymers with tunable thermal and optical properties.

- 5 For example, random amorphous co-polymers with variable refractive indices, glass transition temperatures, and long term thermal stability above 350 degrees C. may be prepared by correct choice of a co-monomer composition. Co-polymers may be prepared by simple melt mixing of variable composition monomer mixtures and heating under appropriate conditions. Using the
10 invention, it is possible under some conditions to provide for precise control of refractive index by the choice of comonomer that is used. Thus, the thermal cyclopolymerization of aromatic trifluorovinyl ether monomers to perfluorocyclobutane polymers and co-polymers affords high temperature, low transmission loss, and variable refractive index for use in optical devices.

15 In one aspect of the invention, optical devices and a process for fabricating optical devices may include an optical waveguide constructed of a core region and a clad region. A lower clad layer is formed on a substrate with a refractive index that is less than that of the core regions.

20 In another aspect of the invention, a core and clad device structure is provided which exhibits a high thermo optic coefficient (-dn/dT) (refractive index change with temperature) for the manufacture of a thermo-optic switch.

Brief Description of the Drawings

A full and enabling disclosure of this invention, including the best mode shown to one of ordinary skill in the art, is set forth in this specification. The
25 following Figures illustrate the invention:

Figure 1 shows the perfluorocyclobutane (PFCB) copolymer composition;

Figure 1A shows the structure of the copolymer that may be formed and used in the practice of the invention;

Figures 2A-2I depict various groups that may be substituted in a poly-aryl ether having perfluorocyclobutyl linkages as shown in Figure 1, wherein the Ar and/or Ar' groups would be substituted onto the linkage shown in Figure 1;

Figure 3 shows a spin coated thickness versus polymer concentration
5 plot;

Figure 4 is a refractive index plot; and

Figure 5 shows a graph of a refractive index versus cure time curve.

Detailed Description of the Invention

Reference now will be made to the embodiments of the invention, one or
10 more examples of which are set forth below. Each example is provided by way
of explanation of the invention, not as a limitation of the invention. In fact, it
will be apparent to those skilled in the art that various modifications and
variations can be made in this invention without departing from the scope or
spirit of the invention. For instance, features illustrated or described as part of
15 one embodiment can be used on another embodiment to yield a still further
embodiment. Thus, it is intended that the present invention cover such
modifications and variations as come within the scope of the appended claims
and their equivalents. Other objects, features and aspects of the present
invention are disclosed in or are obvious from the following detailed description.
20 It is to be understood by one of ordinary skill in the art that the present
discussion is a description of exemplary embodiments only, and is not intended
as limiting the broader aspects of the present invention, which broader aspects
are embodied in the exemplary constructions.

Poly-aryl ethers containing perfluorocyclobutyl (PFCB) linkages are
25 viable candidates for optical and opto-electronic applications due in part to their
wide range of complementary properties. The properties of such compounds
include excellent thermal and thermo-oxidative stability, high T_g, low moisture
absorption, low dielectric constant, variable refractive index, and desirable melt
and solution processability. Furthermore, poly-aryl ethers containing PFCB
30 linkages are among the few known optical polymers to possess low attenuations

in the technologically important 1,550 and 1,300 nm region. And can be precisely index matched to silica optical fibers and exhibit a high thermo-optic coefficient.

- PFCB polymers are prepared from trifluorovinyl aromatic ether monomers via a free radical mediated thermal co-polymerization mechanism as illustrated in Figure 1. A representative example of some of the groups that may be included and used as the Ar and/or Ar' group as shown in Figure 1 are included in Figures 2A-2I.

- A variety of monomers may be used to produce the PFCB polymers. Bis-and tris-functionalized TVE monomers polymerize to form linear or network polymers respectively with unique optical, thermal and mechanical properties. Furthermore, accurate control over the polymer properties may be achieved by co-polymerization.

- It has been surprisingly discovered in the practice of the invention that compositions formed from structures as shown in Figures 1 and 2 may be advantageously used to provide high solids content solutions for coating on substrates in the construction of optical devices, and optical waveguides. High solids content solutions afford the opportunity to build relatively thick films upon the surface of a substrate, which is very desirable in the construction of such devices.

- In the practice of one aspect of the invention, it is possible to copolymerize specific comonomers to provide variable and relatively thick core and clad copolymers for waveguide applications such as switches, modulators, attenuators, and the like. Alternatively, mixtures of homopolymer oligomers could be dissolved in a solvent to provide a similar solution, and then coated and cured. By thick, it is meant having a thickness of at least about 0.6 microns. In some applications, a thickness of 1 micron or more may be achieved in only one coat. In other preferred embodiments of the invention, a thickness of at least about 2 microns may be achieved. Thickness levels from 2-10 microns also can be achieved, and in some aspects of the invention, it may be possible to achieve

thickness levels well over 10 microns, up to and including at least about 50 microns in some particular embodiments of the invention.

The solution processability of the compositions in the practice of the invention is especially favorable. A reactive copolymer is provided which can
5 be dissolved in very low amounts of solvent so that especially thick coatings can be prepared. The films are then thermally cured (i.e. advanced or polymerized) until they comprise three dimensional "network" polymer. The coatings then may be patterned by lithography or reactive ion etching ("RIE") or such as electron beam methods, or micromolding. In general, the network polymer
10 formed is not soluble and will withstand other processing steps, which is another advantageous feature of the invention. The first layer structures may be coated more than once until a core/clad optical device is formed that is multilayered. High solids content of the copolymer solutions, and a suitable copolymer with appropriate properties, are necessary factors to consider in the practice of the
15 invention.

Monomers shown in Figures 2A-2I may be used for optical applications. Bis and tris functionalized TVE monomers polymerize to form linear or network polymers respectively with unique optical, thermal and mechanical properties. Accurate control over polymer properties can be
20 achieved by copolymerization. In addition, methods have been developed that preclude the need for elastomeric negative molds for more cost effective and reproducible prototypes, as further shown below.

PFCB polymers can be prepared neat or in solution at temperatures in excess of 160°C and do not require initiators or catalysts. The polymerization
25 proceeds via a thermally activated cyclodimerization process. Although free-radical mediated, the cyclo-addition polymerization follows classical step growth kinetics. The polymers can be advanced to precisely controlled viscosity, molecular weight and dispersity by simple heating. The bifunctional TVE monomers yield linear polymers with M_n approaching 100,000 whereas
30 trifunctional monomers can be melt advanced to branched pre-networks and

processed prior to final cure. PFCB materials, in general, possess excellent thermal ($T_g=120\text{-}350\text{ }^\circ\text{C}$), mechanical and environmental properties providing long-term stability at about $350\text{ }^\circ\text{C}$.

The step growth kinetics by which PFCB polymers are formed allows for
5 easy control of parameters important to coating technology. Monomer can be
solution advanced at 150 degrees C. in typical solvents known in the art. A
precisely controlled viscosity may be achieved, with correct molecular weight
and polydispersity. The pre-polymer solution may be spin coated, dip coated,
sprayed, or otherwise applied to the substrate by techniques employed in the art
10 of optical device fabrication. Spin coating is known to provide high quality,
defect free films. The film typically is cured by baking under nitrogen at
temperatures ranging from about 235 degrees to about 325 degrees C. for several
hours depending upon the application.

Solids content of solutions of PFCB copolymer employed may be from
15 about 10% to about 70% films or plaques can be prepared by melt-mixing
variable composition monomer mixtures and heating at 150-200 $^\circ\text{C}$ under an
inert atmosphere.

One useful copolymer composition is made by polymerizing the species
shown in Figures 2A and 2B (50 wt% compositions) which has been known to
20 provide a measured $T_g = 220\text{ }^\circ\text{C}$.

Copolymer solutions suitable for spin coating variable thick films may
be prepared in mesitylene solution , yet other solvents such as N-
methylpyrrolidinone and dimethylsulfoxide also can be used due to the wide
ranging and excellent solubility exhibited by trifluorovinyl ether monomers and
25 pre-network copolymer intermediates. For optical device fabrication, many
copolymers studied to date contain the trifunctional monomer shown in Figure
2B so that the advanced intermediate is a branched copolymer of variable
conversion and thus molecular weight and viscosity may be controlled prior to
thermal cure to a cross-linked network.

For example, copolymer solutions in excess of 80 wt % polymer can be obtained (see Table1). This enables the preparation of thick films (on the order of 10 μm) in one spin coat. The ability to spin coat very thick films is a unique feature for high performance PFCB polymers and thereby ensures a
5 homogeneous layer in the final structure versus the multiple spin and cure cycles needed otherwise to achieve the desired thickness.

Figure 3 illustrates one spin curve for polymer solutions formed from copolymers utilizing the two monomer species shown in Figures 2A/2B, ranging from 10-70 weight percent solids at 600 rpm. The range can be tuned precisely
10 with the expected change with spin rate.

Figure 4 illustrates the refractive index (@ 1550 & 653 nm) tunability for two example copolymers as a function of composition. As shown, a wide range of indices can be obtained through choice of composition.

Process parameters can be precisely controlled by choice of comonomer
15 and polymerization time and temperature. After spin coating, the cyclopolymerization is continued by heating under inert atmosphere to the desired second glass transition temperature (T_g). Table 1 illustrates selected conditions and properties for PFCB copolymers targeted for optical applications.

TABLE 1

Condition or Property	Controlled Range
Copolymer solution concentration (mesitylene)	Up to 90 wt %
Single spin coat thickness	Up to 15 μm
Copolymerization temperature / time	140-160 °C / 1 – 10 h
Weight average molecular weight (GPC)	(500 – 50,000)
Molecular weight distribution (GPC)	1.2 – 20 M_w/M_n
Solution viscosity (RMS)	0.02 – 100 Pa s
Cure temperature / time	120 – 350 °C / 0.1 – 3 h
Glass Transition Temperature (DSC T_g)	175 – 350 °C
Patterning Technique	Micromolding or RIE
Loss at 1550 nm	< 0.25 dB/cm
Refractive index at 1550 nm	1.449 – 1.508
Birefringence	< 0.01

The attenuation at 1550 nm for select copolymers is lower than that

- 5 reported in the prior art which used homopolymers of monomer 2B in Fig. 2 instead of copolymers. That is, the use of the monomer in Figure 2B, is lower due to the decrease of aliphatic groups contained only in that particular species (2B) and the intimate compatibility at the core / clad interface for waveguide structures made solely from PFCB copolymers.

10 The evolution and stability of optical properties is also of major concern for processing and performance of polymer integrated optics. The stability of the polymer's optical properties over time and during multiple heat and cool

cycles is currently a significant issue. Initial results using polymers of the invention are shown in Figure 5 where the refractive index is plotted as a function of cure time at 200 °C for the monomer of Figure 2B. As shown, the refractive index changes dramatically during cure yet quickly stabilizes in less than 2 hours. Further, the refractive index is completely stable after multiple heat cure cycles and continued heating at 200 °C.

The copolymers of the invention exhibit relatively low loss in the telecommunications wavelength at which they are employed. That is, a low optical loss is observed at about 1550 nm. Loss levels of less than about 0.3, and sometimes less than about 0.1 dB/cm, may be achieved using the compositions and or more, with greater thickness film layers being achieved using greater solids content percentages. Solvents used may be any of the commonly known solvents, including for example, mesitylene. Spin coating is performed by either pouring the solution (less than 10 ml) upon a static silicon wafer (secured by vacuum on the spin coater chuck) and spinning at the desired speed and time (electronically controlled), or pouring the solution on the spinning wager.

General Procedure Experimental Details

Monomers were prepared from their corresponding phenolic precursors. ¹H NMR data were obtained on a Bruker AC-300 spectrometer. ¹⁹F NMR 188 MHz data were obtained on a Bruker AC-200 spectrometer. Functional group (TVE) conversion data (Table 2) was determined by integration of the TVE fluorine signals and PFCB fluorine signals. Chloroform-d was used as solvent, and chemical shifts reported are internally referenced to tetramethylsilane (0 ppm), CDCl₃ (77 ppm), and CFCl₃ (0 ppm) for ¹H, ¹³C, and ¹⁹F nuclei, respectively. Infrared analyses were performed on neat oils or free standing films using a Nicolet 550 Magna FTIR spectrophotometer. IR data used for polymerization kinetics was obtained in reflectance mode on an ASI React IR System. Gas chromatography/mass spectrometry (GC/MS) data were obtained from a Varian Saturn GC/MS. Gel permeation chromatography (GPC) data were collected in

THF using a Waters 2690 Alliance System with refractive index detection at 35 °C, and equipped with two consecutive Polymer Labs PLGel 5mm Mixed-D and Mixed-E columns. Retention times were calibrated against Polymer Labs Easical PS-2 polystyrene standards. The copolymer solution viscosity was measured with a TA AR1000-N Rheometer. A Geometry of cone-plate was used with a diameter of 40 mm, the 2 degree angle and 43 µm truncation. The frequency dependence of viscosity was measured within the frequency range of 3.0 to 300.0 rad/s at room temperature. The viscosity data reported here is read at the shear rate of 100.0 rad/s. DSC data were obtained from a Mettler-Toledo 820 System under a nitrogen atmosphere at a scan-rate of 10°C/min.

Table 2 Selected Copolymer Data

Ex.	Monomer (g)	Monomer (g)	wt.% polymer in mesitylene	Polym. time (h) at 150/160 °C	Mn	Mw	Mw/Mn	Olefin conv. (%)	Viscosity (Pa.s)	*Refractive index @1550 nm	*Glass Transition, Tg (°C)
1	2B(175)	0	75	1.45	2444	7033	2.89	34.8	0.1	1.4883	350
2	2B(40)	2A(40)	80	2.5	589	1212	2.06	37.3	0.133	1.5036	225
3	2B(17.38)	2C(7.78)	50	8	1859	4656	2.5	44.2	6.8	1.4785	221
4	2B(17.38)	2C(7.78)	75	5	3943	9709	2.45	52.2	1.575	1.4801	220
5	2B(50)	2A(50)	70	2.15	661	1178	1.78	37	1.1	1.5008	224
6	2B(160)	2A(40)	75	3.2	2508	4846	1.93	40	0.1	1.4892	285
7	2B(80)	2A(20)	70	1.45	899	2008	2.34	32	0.085	1.4892	286

* After cure at 200 °C for 2 h.

Copolymer solutions were prepared by dissolving one or more monomers as shown in Figs. 2A-2B (50-80 wt %) in freshly distilled and filtered mesitylene (5 L flask, ca. 3 L polymer solution). The solution was stirred mechanically and degassed for 30 minutes with a dry nitrogen purge. The purge was stopped and a positive pressure of nitrogen was applied and the reaction flask was heated to 150 °C and stirred for several hours to form the copolymers.

The copolymer films were applied to silicon wafers using a KW-4A Spincoater (Chemate Technology, Inc.) using standard spin coating techniques. Filtered polymer solutions were coated from 600 to 2000 rpm for about 60 seconds, and then cured in an oven 200 °C under nitrogen for 2 hours.

Alternatively, films could be hot plate cured under nitrogen, in other applications of the invention. Also, the temperature could potentially be increased, or decreased, depending upon the particular application and copolymer employed.

- 5 Film thickness was measured by a profilometer, Alphastep 200, which is manufactured by Tencor Company. Refractive index data was obtained from a Metricon Model 2010 Prism Coupler System from Metricon Company. Birefringence was calculated as the difference between the TE mode and the TM mode ($\eta_{TE} - \eta_{TM}$). Absorption spectra were measured using a Perkin Elmer
10 Lambda 900 UV/Vis/NIR spectrophotometer. Spectra were taken over the wavelength range 3200-200 nm at a scan rate of 120 nm/minute.

15 **Example 1: Homopolymer of 2B (75 wt% in mesitylene)**

To a three neck round bottom flask equipped with nitrogen inlet, reflux condenser and mechanical stir was added 175 g of pure monomer **2B** and 58.3 g of filtered mesitylene. The solution was degassed as described in the general procedures and experimental details above. The clear solution was then heated
20 to 150-160 °C for 1.45 h and allowed to cool. The resulting homopolymer exhibited a olefin conversion of 34.8 %, a number average molecular weight (Mn) of 2,444 and a weight average molecular weight (Mw) of 7033, and a molecular weight distribution of 2.89 (See Table 2). The polymer solution was spin coated and cured as described above and gave a thermoset film with
25 refractive index of 1.4883 (at 1550 nm) and a glass transition temperature of 350 °C.

Example 2: Copolymer of 2B and 2A (50:50 ratio, 80 wt% in mesitylene)

To a three neck round bottom flask equipped with nitrogen inlet, reflux
30 condenser and mechanical stir was added 40 g of pure monomer **2B**, 40 g of pure monomer **2A**, and 10 g of filtered mesitylene. The solution was degassed as described in the general procedures and experimental details above. The clear

solution was then heated to 150-160 °C for 2.5 h and allowed to cool. The resulting copolymer exhibited a olefin conversion of 37.3 %, a number average molecular weight (Mn) of 589 and a weight average molecular weight (Mw) of 1212, and a molecular weight distribution of 2.06 (See Table 2). The polymer solution was spin
5 coated and cured as described above and gave a thermoset film with refractive index of 1.5036 (at 1550 nm) and a glass transition temperature of 225 °C.

Example 3: Copolymer of 2B and 2C (70:30 ratio, 50 wt% in mesitylene)

To a three neck round bottom flask equipped with nitrogen inlet, reflux
10 condenser and mechanical stir was added 17.4 g of pure monomer **2B**, 7.8 g of pure monomer **2C**, and 25 g of filtered mesitylene. The solution was degassed as described in the general procedures and experimental details above. The clear solution was then heated to 150-160 °C for 8 h and allowed to cool. The resulting copolymer exhibited a olefin conversion of 44.2 %, a number average molecular
15 weight (Mn) of 1859 and a weight average molecular weight (Mw) of 4656, and a molecular weight distribution of 2.5 (See Table 2). The polymer solution was spin coated and cured as described above and gave a thermoset film with refractive index of 1.4785 (at 1550 nm) and a glass transition temperature of 221 °C.

Example 4: Copolymer of 2B and 2C (70:30 ratio, 75 wt% in mesitylene)

To a three neck round bottom flask equipped with nitrogen inlet, reflux condenser and mechanical stir was added 17.4 g of pure monomer **2B**, 7.8 g of pure monomer **2C**, and 8 g of filtered mesitylene. The solution was degassed as described in the general procedures and experimental details above. The clear solution was then heated to 150-160 °C for 5 h and allowed to cool. The resulting copolymer exhibited a olefin conversion of 52.2 %, a number average molecular weight (Mn) of 3943 and a weight average molecular weight (Mw) of 9709, and a molecular weight distribution of 2.45 (See Table 2). The polymer solution was spin coated and cured as described above and gave a thermoset film with refractive
30 index of 1.4801 (at 1550 nm) and a glass transition temperature of 220 °C.

Example 5: Copolymer of 2B and 2A (50:50 ratio, 70 wt% in mesitylene)

To a three neck round bottom flask equipped with nitrogen inlet, reflux condenser and mechanical stir was added 50 g of pure monomer **2B**, 50 g of pure monomer **2A**, and 43 g of filtered mesitylene. The solution was degassed as described in the general procedures and experimental details above. The clear solution was then heated to 150-160 °C for 2.15 h and allowed to cool. The resulting copolymer exhibited a olefin conversion of 37 %, a number average molecular weight (Mn) of 661 and a weight average molecular weight (Mw) of 1178, and a molecular weight distribution of 1.78 (See Table 2). The polymer solution was spin coated and cured as described above and gave a thermoset film with refractive index of 1.5008 (at 1550 nm) and a glass transition temperature of 224 °C.

Example 6: Copolymer of 2B and 2A (80:20 ratio, 75 wt% in mesitylene)

To a three neck round bottom flask equipped with nitrogen inlet, reflux condenser and mechanical stir was added 160 g of pure monomer **2B**, 40 g of pure monomer **2A**, and 67 g of filtered mesitylene. The solution was degassed as described in the general procedures and experimental details above. The clear solution was then heated to 150-160 °C for 3.2 h and allowed to cool. The resulting copolymer exhibited a olefin conversion of 40 %, a number average molecular weight (Mn) of 2508 and a weight average molecular weight (Mw) of 4846, and a molecular weight distribution of 1.93 (See Table 2). The polymer solution was spin coated and cured as described above and gave a thermoset film with refractive index of 1.4892 (at 1550 nm) and a glass transition temperature of 285 °C.

Example 7: Copolymer of 2B and 2A (80:20 ratio, 70 wt% in mesitylene)

To a three neck round bottom flask equipped with nitrogen inlet, reflux condenser and mechanical stir was added 80 g of pure monomer **2B**, 40 g of pure monomer **2A**, and 20 g of filtered mesitylene. The solution was degassed as described in the general procedures and experimental details above. The clear solution was then heated to 150-160 °C for 1.45 h and allowed to cool. The

resulting copolymer exhibited a olefin conversion of 32 %, a number average molecular weight (Mn) of 899 and a weight average molecular weight (Mw) of 2008, and a molecular weight distribution of 2.34 (See Table 2). The polymer solution was spin coated and cured as described above and gave a thermoset film
5 with refractive index of 1.4892 (at 1550 nm) and a glass transition temperature of 286 °C.

Other solvents besides mesitylene could be used in the practice of the invention, such as other high boiling hydrocarbons, N-methylpyrrolidinone, dimethylsulfoxide, and others. For optical device fabrication, most copolymer
10 intermediates enable molecular weight and viscosity control prior to thermal cure to a cross-linked network.

It is understood by one of ordinary skill in the art that the present discussion is a description of exemplary embodiments only, and is not intended as limiting the broader aspects of the present invention, which broader aspects
15 are embodied in the exemplary constructions. The species shown in Figures 2A-2I could be employed in numerous different combinations in the practice of the invention, at various ratios and the like. The invention is shown by example in the appended claims.